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(54) Title: AMIDE DERIVATIVES AS 5HT1D RECEPTOR ANTAGONISTS

(57) Abstract

Novel amide derivatives, processes for their preparation, pharmaceutical compositions containing them and their use as medicaments are disclosed.

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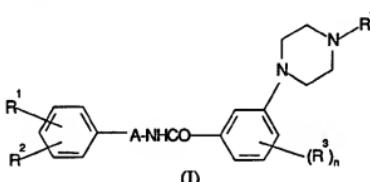
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Amide derivatives as 5HT_{1D} receptor antagonists

The present invention relates to novel amide derivatives, processes for their preparation, and pharmaceutical compositions containing them.

5 EPA 0 533 266/7/8 disclose a series of benzanilide derivatives which are said to possess 5HT_{1D} receptor antagonist activity. These compounds are alleged to be of use in the treatment of various CNS disorders.

10 A structurally distinct class of compounds have now been discovered and have been found to exhibit 5HT_{1D} antagonist activity. In a first aspect, the present invention therefore provides a compound of formula (I) or a salt thereof:



15 in which

R¹ is optionally substituted phenyl or an optionally substituted 5 to 7-membered heterocyclic ring containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulphur or R¹ is hydrogen, halogen, C₁-6alkyl, C₁-6cycloalkyl, C₁-6alkoxy, hydroxyC₁-6alkyl, acyl, nitro, trifluoromethyl, cyano, SR⁵, SOR⁵, SO₂R⁵, SO₂NR⁵R⁶, CO₂R⁵, CONR⁵R⁶,

20 CONR⁵(CH₂)_pCO₂R⁶, NR⁵R⁶, NR⁵CO₂R⁶, CR⁵=NOR⁶,

where R⁵ and R⁶ are independently hydrogen or C₁-6alkyl and p is 1 to 3;

R² is hydrogen, halogen, C₁-6alkyl, C₁-6alkoxy, hydroxyC₁-6alkyl, acyl, nitro, trifluoromethyl, cyano, SR⁵, SOR⁵, SO₂R⁵, SO₂NR⁵R⁶, CO₂R⁵, CONR⁵R⁶, CONR⁵(CH₂)_pCO₂R⁶, NR⁵R⁶, NR⁵CO₂R⁶, CR⁵=NOR⁶,

25 where R⁵, R⁶ and p are as defined for R¹;

R³ is hydrogen, halogen, hydroxy, C₁-6alkyl or C₁-6alkoxy;

R⁴ is hydrogen or C₁-6alkyl;

A is a bond or an acyclic hydrocarbon chain having 1 to 6 carbon atoms; and

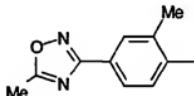
n is 1 or 2.

30 The group R¹ can be an aromatic or saturated heterocyclic ring. When R¹ is an aromatic heterocyclic ring, examples of such rings include thieryl, furyl, pyrrolyl, triazolyl, diazolyl, isothiazolyl, oxadiazolyl, isoxazolyl, thiadiazolyl, pyrimidyl and pyrazinyl. When R¹ is a saturated ring examples include piperidine, morpholine and

piperazine rings. The group R¹ can be linked to the remainder of the molecule via a carbon atom or, when present, a nitrogen atom.

Preferably the group R¹ is attached to the 4-position of the phenyl ring, that is to say, para to the amide group. Optional substituents for R¹ heterocyclic rings include 5 halogen, C₁-6alkyl, C₁-6alkoxy, hydroxy, cyano, nitro, amino, CO₂R⁵ or CONR⁵R⁶ where R⁵ and R⁶ are independently hydrogen or C₁-6alkyl. R¹ phenyl groups can be mono or di-substituted. Optional substituents for R¹ phenyl groups include halogen, C₁-6alkyl, C₁-6alkoxy, hydroxy, cyano, nitro, amino, CO₂R⁵ or CONR⁵R⁶ where R⁵ and R⁶ are independently hydrogen or C₁-6alkyl as well as 5 to 7-membered heterocyclic 10 rings containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulphur. These heterocyclic rings can themselves be substituted, for example by C₁-6alkyl.

Preferably R¹ is halogen, cyclohexyl, optionally substituted pyridyl or optionally substituted phenyl. More preferably R¹ is phenyl disubstituted by a C₁-6alkyl group and a 1,2,4-oxadiazol-3-yl group, in particular disubstituted by a methyl and a 5-methyl-1,2,4-15 oxadiazol-3-yl group. Most preferably R¹ is a group of formula:



Suitably R² is hydrogen, halogen, C₁-6alkyl, C₁-6alkoxy, hydroxyC₁-6alkyl, 20 acyl, nitro, trifluoromethyl, cyano, SR⁵, SOR⁵, SO₂R⁵, SO₂NR⁵R⁶, CO₂R⁵, CONR⁵R⁶, CONR⁵(CH₂)_pCO₂R⁶, NR⁵R⁶, NR⁵CO₂R⁶, CR⁵=NOR⁶ where R⁵ and R⁶ are independently hydrogen or C₁-6alkyl and p is 1 to 3. Preferably R² is hydrogen or C₁-6alkyl, for example methyl.

Suitably R³ is hydrogen, halogen, hydroxy, C₁-6alkyl or C₁-6alkoxy. Preferably 25 R³ is C₁-6alkoxy such as methoxy.

Preferably n is 1.

Suitably R⁴ is hydrogen or C₁-6alkyl. Preferably R⁴ is C₁-6alkyl such as methyl.

For the avoidance of doubt, the term 'chain of 1 to 6 carbon atoms' means carbon atoms extending in a branched or unbranched chain between the phenyl group and the 30 amide group. The hydrocarbon chain can be an alkylene chain, for example methylene or ethylene, or A can contain alkene or alkyne groups. For example, the group A can be methylene or ethylene. Preferably A is a bond.

C₁-6alkyl groups, whether alone or as part of another group, may be straight chain or branched.

Particularly preferred compounds include:

- 4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromo-3-methylphenyl)benzamide,
- 4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromophenyl)benzamide,
- 4-methoxy-3-(4-methyl-1-piperazinyl)-N-[3-methyl-4-(4-pyridyl)phenyl]benzamide,
- 5 4-methoxy-3-(4-methyl-1-piperazinyl)-N-[2'-methyl-4'-(5-methyl-1,2,4-oxadiazol-3-yl)(1,1'-biphenyl)]benzamide, or
- 4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-cyclohexylphenyl)benzamide, or pharmaceutically acceptable salts thereof.

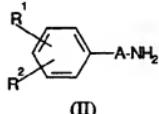
Preferred salts of the compounds of formula (I) are pharmaceutically acceptable salts. These include acid addition salts such as hydrochlorides, hydrobromides, phosphates, acetates, fumarates, maleates, tartrates, citrates, oxalates, methane sulphonates and p-toluenesulphonates.

Certain compounds of formula (I) are capable of existing in stereoisomeric forms. It will be understood that the invention encompasses all geometric and optical isomers of the compounds of formula (I) and the mixtures thereof including racemates. Tautomers of compounds of formula (I) and mixtures thereof also form part of the invention.

In a further aspect the present invention provides a process for the preparation of a compound of formula (I) which comprises:

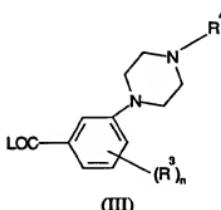
(a) reaction of a compound of formula (II):

20



25

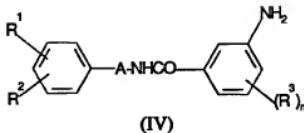
in which R¹, R² and A are as defined in formula (I), with a compound of formula (III):



30

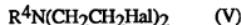
in which R³, R⁴ and n are as defined in formula (I) and L is a leaving group; or

(b) reaction of a compound of formula (IV):



5

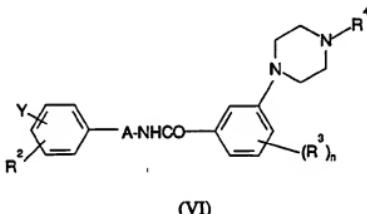
in which R^1 , R^2 , R^3 , A and n are as defined in formula (I) with a compound of formula (V):



10

in which R^4 is as defined in formula (I) and Hal is halogen, or

(c) reaction of a compound of formula (VI):



15

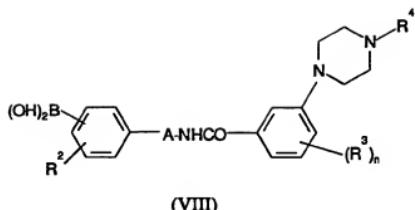
in which R^2 , R^3 , R^4 , A and n are as defined in formula (I) and Y is halogen or a group -
20 OSO_2CF_3 with a compound of formula (VII):



25

in which R^1 is as defined in formula (I), or

(d) reaction of a compound of formula (VIII):



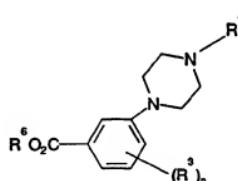
- 5 in which R^2 , R^3 , R^4 , A and n are as defined in formula (I) with a compound of formula (IX):



- 10 (IX)

in which R^1 is as defined in formula (I) and Y is as defined in formula (VI), or

- (e) reaction of a compound of formula (II) as defined above with a
15 compound of formula (X):



- 20 in which R^3 , R^4 and n are as defined in formula (I) and R^6 is C_{1-4} -alkyl in the presence of a trialkylaluminium reagent;

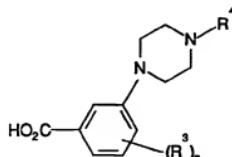
and optionally after any of the above processes:

- 25 • converting a compound of formula (I) into another compound of formula (I)
• forming a pharmaceutically acceptable salt.

Suitable activated carboxylic acid derivatives of formula (III) include acyl halides and acid anhydrides. Activated compounds of formula (III) can also be prepared by reaction of the corresponding carboxylic acid with a coupling reagent such as

carbonyldiimidazole, dicyclohexylcarbodiimide or diphenylphosphorylazole. Preferably the group L is halo, particularly chloro.

- A compound of formula (II) is typically reacted with a compound of formula (III) in an inert organic solvent such as DMF, THF or dichloromethane at ambient or elevated 5 temperature in the presence of a base such as triethylamine or pyridine.
Compounds of formula (III) can be prepared from a compound of formula (XI):



10 (XI)

in which R^3 , R^4 and n are as defined in formula (I) using standard procedures. For example acid chlorides can be prepared by reaction with phosphorous pentachloride, 15 oxalyl chloride or thionyl chloride. Acid anhydrides can be prepared by reaction with a suitable acid anhydride, for example trifluoroacetic anhydride.

Reaction of a compound of formula (IV) with a compound of formula (V) is suitably carried out in an alcohol or nitrile solvent with an optional base or, alternatively, in a non-polar solvent such as chlorobenzene in the absence of base. Suitably, the reactions are carried out at ambient or elevated temperature, preferably at the reflux 20 temperature of the reaction mixture.

Reaction of compounds of formula (VI) and (VII) and reaction of compounds of formulae (VIII) and (IX) can be carried out in the presence of a transition metal catalyst such as $\text{Pd}(\text{PPh}_3)_4$ in a solvent such as an ether in the presence of a base such as an alkali metal carbonate or bicarbonate, for example sodium carbonate or bicarbonate, at ambient 25 or elevated temperature.

Compounds of formula (II) can be reacted with compounds of formula (X) in an organic solvent such as toluene in an inert atmosphere in the presence of a trialkylaluminium reagent, for example trimethylaluminium.

Intermediate compounds of formulae (II), (IV), (V), (VI), (VII), (VIII), (IX), (X) 30 and (XI) are either commercially available or can be prepared using standard procedures. Certain intermediate compounds of formulae (II), (IV), (V), (VI), (VII), (VIII), (IX), (X) and (XI) are novel and form a further aspect of the invention.

It will be appreciated to those skilled in the art that it may be necessary to protect certain reactive substituents during some of the above procedures, for example when the

group R⁴ is a hydrogen atom. Standard protection and deprotection techniques can be used. For example, primary amines can be protected as phthalimide, benzyl, benzyloxycarbonyl or trityl derivatives. These groups can be removed by conventional procedures.

- 5 Carboxylic acid groups can be protected as esters. Aldehyde or ketone groups can be protected as acetals, ketals, thioacetals or thioketals. Deprotection is achieved using standard conditions.

- 10 Compounds of formula (I) can be converted into further compounds of formula (I) using standard techniques. For example, in the case wherein R⁴ is hydrogen, it is possible to introduce a C₁-alkyl group by conventional alkylation using 1 molar equivalent of a C₁-alkyl halide and 1 molar equivalent of a suitable base in an inert solvent. Compounds of formula (I) in which R¹ or R² are acid groups can be esterified using normal procedures.

- 15 5HT_{1D} Antagonists, and in particular the compounds of the present invention, are expected to be of use in the treatment of CNS disorders such as mood disorders, including depression, seasonal effective disorder and dysthymia; anxiety disorders, including generalised anxiety, panic disorder, agoraphobia, social phobia, obsessive compulsive disorder and post-traumatic stress disorder; memory disorders, including dementia, amnestic disorders and age-associated memory impairment; and disorders of eating behaviours, including anorexia nervosa and bulimia nervosa. Other CNS disorders include Parkinson's disease, dementia in Parkinson's disease, neuroleptic-induced Parkinsonism and tardive dyskinésias, as well as other psychiatric disorders.

- 20 25 5HT_{1D} Antagonists, and in particular compounds of the present invention, may also be of use in the treatment of endocrine disorders such as hyperprolactinaemia, in the treatment of vasospasm (particularly in the cerebral vasculature) and hypertension, as well as disorders in the gastrointestinal tract where changes in motility and secretion are involved. They may also be of use in the treatment of sexual dysfunction.

- Therefore, the present invention, provides a compound of general formula (I) or a physiologically acceptable salt or solvate thereof for use in therapy.
- 30 The present invention also provides a compound of general formula (I) or a physiologically acceptable salt or solvate thereof for use in the treatment of the aforementioned disorders.

- In another aspect the invention provides the use of a compound of general formula (I) or a pharmaceutically acceptable salt or solvate thereof for the manufacture of a medicament for the treatment of the aforementioned disorders.

35 In a further aspect the invention provides a method of treating the aforementioned disorders which comprises administering an effective amount to a patient in need of such

treatment of a compound of general formula (I) or a pharmaceutically acceptable salt or solvate thereof.

In particular the invention provides a compound of general formula (I) or a physiologically acceptable salt or solvate thereof for use in the treatment or prophylaxis of depression.

It will be appreciated by those skilled in the art that the compounds according to the invention may advantageously be used in conjunction with one or more other therapeutic agents, for instance, different antidepressant agents.

The present invention also provides a pharmaceutical composition, which 10 comprises a compound of formula (I) or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

A pharmaceutical composition of the invention, which may be prepared by admixture, suitably at ambient temperature and atmospheric pressure, is usually adapted for oral, parenteral or rectal administration and, as such, may be in the form of tablets, 15 capsules, oral liquid preparations, powders, granules, lozenges, reconstitutable powders, injectable or infusible solutions or suspensions or suppositories. Orally administrable compositions are generally preferred.

Tablets and capsules for oral administration may be in unit dose form, and may contain conventional excipients, such as binding agents, fillers, tabletting lubricants, 20 disintegrants and acceptable wetting agents. The tablets may be coated according to methods well known in normal pharmaceutical practice.

Oral liquid preparations may be in the form of, for example, aqueous or oily suspension, solutions, emulsions, syrups or elixirs, or may be in the form of a dry product for reconstitution with water or other suitable vehicle before use. Such liquid preparations 25 may contain conventional additives such as suspending agents, emulsifying agents, non-aqueous vehicles (which may include edible oils), preservatives, and, if desired, conventional flavourings or colorants.

For parenteral administration, fluid unit dosage forms are prepared utilising a compound of the invention or pharmaceutically acceptable salt thereof and a sterile 30 vehicle. The compound, depending on the vehicle and concentration used, can be either suspended or dissolved in the vehicle. In preparing solutions, the compound can be dissolved for injection and filter sterilised before filling into a suitable vial or ampoule and sealing. Advantageously, adjuvants such as a local anaesthetic, preservatives and buffering agents are dissolved in the vehicle. To enhance the stability, the composition 35 can be frozen after filling into the vial and the water removed under vacuum. Parenteral suspensions are prepared in substantially the same manner, except that the compound is suspended in the vehicle instead of being dissolved, and sterilisation cannot be

accomplished by filtration. The compound can be sterilised by exposure to ethylene oxide before suspension in a sterile vehicle. Advantageously, a surfactant or wetting agent is included in the composition to facilitate uniform distribution of the compound.

5 The composition may contain from 0.1% to 99% by weight, preferably from 10 to 60% by weight, of the active material, depending on the method of administration.

The dose of the compound used in the treatment of the aforementioned disorders will vary in the usual way with the seriousness of the disorders, the weight of the sufferer, and other similar factors. However, as a general guide suitable unit doses may be 0.05 to 1000 mg, more suitably 1.0 to 200 mg, and such unit doses may be administered more 10 than once a day, for example two or three a day. Such therapy may extend for a number of weeks or months.

When administered in accordance with the invention, no unacceptable toxicological effects are expected with the compounds of the invention.

15 The following Examples illustrate the preparation of pharmacologically active compounds of the invention.

Description 1**Methyl 4-methoxy-3-(4-methyl-1-piperazinyl)benzoate**

Methyl 4-hydroxy-3-nitrobenzoate (5.00g, 0.025mol), was dissolved in acetone (100 ml),
5 and treated with anhydrous potassium carbonate (6.91g, 0.050mol), followed by iodomethane (1.71 ml, 0.0275mol). The mixture was then heated to reflux with stirring. After 16h, more iodomethane (1.71 ml, 0.0275mol) was added and reflux was maintained for a further 3h. The reaction mixture was then filtered and the filtrate evaporated under reduced pressure to give an orange solid, which was dried *in vacuo*. The solid was then
10 suspended in ethanol (200 ml), and was hydrogenated at atmospheric pressure in the presence of 5% PdC catalyst (0.5g). After 16h, the reaction mixture was filtered through kieselguhr and evaporated under reduced pressure to give a pale yellow oily solid, which was dried *in vacuo*. The oily solid was dissolved in chlorobenzene (50 ml), and mechloroethamine hydrochloride (14.44g, 0.075mol) was added. The reaction mixture was
15 then heated to reflux under argon. After 30h, the reaction mixture was allowed to cool, and was left for a further 24h at room temperature before being evaporated under reduced pressure and partitioned between potassium carbonate solution and dichloromethane. The aqueous layer was then extracted with dichloromethane, and the combined organic layers were dried (Na_2SO_4) and evaporated under reduced pressure to give a brown oily solid,
20 which was dried *in vacuo*, and purified by silica gel chromatography (5% MeOH/ CH_2Cl_2) to give the title compound (0.740g, 11%) as a brown oil.

^1H NMR (250 MHz, CDCl_3), δ 7.75 (dd, 1H), 7.61 (s, 1H), 6.86 (d, 1 H), 3.93 (s, 3H),
3.87 (s, 3H), 3.14 (br s, 4H), 2.65 (br s, 4H), 2.38 (s, 3H).

25

Example 1**4-Methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromo-3-methylphenyl)benzamide**

30 4-Bromo-3-methylaniline (0.2g, 1.08 mmol) was dissolved in dry toluene (10 ml) and treated with trimethylaluminium (2.0 M in toluene) (2.16ml, 4.32 mmol) with stirring under argon. After 0.25h, a solution of the product from description 1 (0.285g, 1.08mmol) in toluene (5 ml) was added. The mixture was then stirred at room temperature for 2h, before being heated to 80°C. After 2 h at 80°C, the reaction mixture was allowed to cool
35 and was treated with dilute HCl, until no more effervescence took place. The reaction mixture was then partitioned between dichloromethane and water. The aqueous layer was then treated with aq. sodium bicarbonate until basic and then extracted with

dichloromethane (2X). The combined organic layers were then dried (Na_2SO_4), and evaporated under reduced pressure to give a pale brown solid. The solid was the purified by silica-gel chromatography (6% $\text{MeOH}/\text{CH}_2\text{Cl}_2$) as eluant, to give the title compound as a cream coloured foam (0.210g, 47%), which was converted to its oxalate salt.

5

m.pt 144-145°C

10 ^1H NMR (200MHz, CDCl_3 -free base- δ) 7.92 (s, 1H), 7.62 (d, 1H), 7.48 (m, 3H), 7.35 (dd, 1H), 6.92 (d, 1H), 3.94 (s, 3H), 3.19 (br s, 4H), 2.73 (br s, 4H), 2.43 (s, 3H), 2.39 (s, 3H).

Example 2

4-Methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromophenyl)benzamide

15 4-Bromoaniline (0.260g, 1.51 mmol) was dissolved in toluene (10 ml), and was treated with trimethylaluminium (2.0M in toluene) (2.30 ml, 4.53 mmol) with stirring under argon. After 0.25h, a solution of the product from description 1 (0.400g, 1.51 mmol) in toluene (10 ml) was added. The mixture was then heated to 80°C. After 2h, the reaction mixture was allowed to cool, and was poured into a slurry of silica-gel (~10g) in dichloromethane (40 ml). The mixture was then stirred carefully until effervescence ceased and filtered. The filter pad was then washed with 20% $\text{MeOH}/\text{CH}_2\text{Cl}_2$ (400ml) and the filtrate was evaporated under reduced pressure to give a brown oil, which was purified by silica-gel chromatography (10% $\text{MeOH}/\text{CH}_2\text{Cl}_2$ as eluant) to give the title compound as a colourless oil (0.475g, 78%), that crystallised on standing.

25

^1H NMR (250 MHz, CDCl_3) δ 7.88 (s, 1H), 7.60-7.45 (m, 6H), 6.88 (d, 1H), 3.93 (s, 3H), 3.15 (br s, 4H), 2.65 (br s, 4H), 2.38 (s, 3H)

25

Example 3

4-Methoxy-3-(4-methyl-1-piperazinyl)-N-[3-methyl-4-(4-pyridyl)phenyl]benzamide

30 The product from example 1 (0.150g, 0.360mmol), 4-pyridylboronic acid (0.044g, 0.360mmol), and tetrakis(triphenylphosphine)palladium(0) (0.025g) were dissolved in DME (8 ml), and sodium carbonate (0.212g, 2.00mmol) in water (2 ml) was added. The mixture was then heated to reflux under argon. After 20 h, further amounts of palladium catalyst (0.025g), and 4-pyridylboronic acid (0.030g, 0.244mmol) were added. After reflux was continued for a further 1h, the reaction mixture was allowed to cool, and was

partitioned between dichloromethane and water. The aqueous layer was then extracted with dichloromethane (1X), and the combined organic layers were dried (Na_2SO_4) and evaporated under reduced pressure to give a brown oil, which was purified by silica-gel chromatography (10% MeOH/ CH_2Cl_2 as eluant) to give the title compound as a pale yellow oil (0.116g, 77%), which was crystallised from 60-80 petrol/diethyl ether.

m.pt 110-111°C

1 H NMR (270 MHz, CDCl_3), δ 8.62 (dd, 2H), 7.90 (s, 1H), 7.65-7.40 (m, 4H), 7.25 (m, 3H), 6.90 (d, 1H), 3.93 (s, 3H), 3.18 (br s, 4H), 2.65 (br s, 4H), 2.40 (s, 3H), 2.30 (s, 3H).

Example 4

4-Methoxy-3-(4-methyl-1-piperazinyl)-N-[2'-methyl-4'-(5-methyl-1,2,4-oxadiazol-3-yl) (1,1'-biphenyl)]benzamide

n-Butyllithium (1.6M in hexanes) (8.36 ml, 0.013 mol) was added dropwise at -90 to -100°C to a stirred solution of the product from Example 1 (0.450g, 1.11 mmol) in dry THF (20 ml) over 20 minutes. The reaction mixture was kept at -90°C for 0.5h, before being allowed to warm to -78°C, and was kept at -78°C for a further 1h, before being allowed to warm to room temperature. The reaction mixture was then stirred at room temperature overnight, before being treated with water (5 ml). After a further 2h stirring at room temperature, the reaction mixture was evaporated under reduced pressure and dried *in vacuo*. The resultant yellow solid was then purified by silica-gel chromatography (10% MeOH/ CH_2Cl_2) to give the derived phenylboronic acid (0.090g) as an off-white solid.

The phenylboronic acid (0.075g, 0.203mmol) was then dissolved in DME (5 ml) and treated with 3-(4-Bromo-3-methylphenyl)-5-methyl-1,2,4-oxadiazole (0.051g, 0.203mmol) (E.P. O 533 268 A1). A solution of sodium carbonate (0.065g, 0.609mmol) in water (5 ml) was added. The mixture was then flushed with argon and $\text{Pd}(\text{PPh}_3)_4$ (10 mg) was added. The mixture was heated to reflux under argon. After 20h, the reaction mixture was allowed to cool, and was partitioned between CH_2Cl_2 and water. The aqueous layer was then extracted with CH_2Cl_2 and the combined organic layers were dried (Na_2SO_4) and evaporated under reduced pressure to give a brown oil which was purified by prep. t.l.c. (5-10% MeOH/ CH_2Cl_2 as eluant) to give the title compound as a pale yellow oil (0.029g), which was converted to its oxalate salt.

m.pt 124-125°C (oxalate salt)

¹H NMR (250 MHz, CDCl₃)-free base- δ 7.97 (m, 2H), 7.72 (d, 2H), 7.55 (d, 2H), 7.38 (m, 3H), 7.27 (s, 1H), 6.91 (d, 1H), 3.94 (s, 3H), 3.19 (br s, 4H), 2.73 (br s, 7H), 2.40 (s, 3H), 2.36 (s, 3H).

5 **Example 5**

4-Methoxy-3-(4-methyl-1-piperazinyl)-N-(4-cyclohexylphenyl)benzamide

The title compound was prepared from 4-cyclohexylaniline (0.19g, 1.1mmol) and the product from description 1 (0.29g, 1.1mmol) using the method described for example 2.

10 Yield 0.13g, 29%.

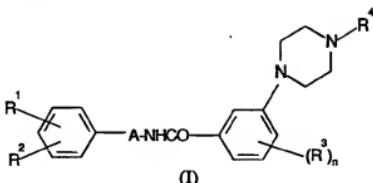
m.pt 146-148°C

¹H NMR (250 MHz, CDCl₃) δ 7.71 (s, 1H), 7.59-7.36 (m, 4H), 7.20 (d, 2H), 6.90 (d,

15 1H), 3.92 (s, 3H), 3.12 (br s, 4H), 2.75-2.41 (m, 5H), 2.35 (s, 3H), 1.98-1.54 (m, 5H),
1.51-1.11 (m, 5H)

CLAIMS:

1. A compound of formula (I) or a salt thereof:



5

in which

R¹ is optionally substituted phenyl or an optionally substituted 5 to 7-membered

10 heterocyclic ring containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulphur or R¹ is hydrogen, halogen, C₁-6alkyl, C₁-6cycloalkyl, C₁-6alkoxy, hydroxyC₁-6alkyl, acyl, nitro, trifluoromethyl, cyano, SR⁵, SOR⁵, SO₂R⁵, SO₂NR⁵R⁶, CO₂R⁵, CONR⁵R⁶, CONR⁵(CH₂)_pCO₂R⁶, NR⁵R⁶, NR⁵CO₂R⁶, CR⁵=NOR⁶,

where R⁵ and R⁶ are independently hydrogen or C₁-6alkyl and p is 1 to 3;

15 R² is hydrogen, halogen, C₁-6alkyl, C₁-6alkoxy, hydroxyC₁-6alkyl, acyl, nitro, trifluoromethyl, cyano, SR⁵, SOR⁵, SO₂R⁵, SO₂NR⁵R⁶, CO₂R⁵, CONR⁵R⁶, CONR⁵(CH₂)_pCO₂R⁶, NR⁵R⁶, NR⁵CO₂R⁶, CR⁵=NOR⁶,

where R⁵, R⁶ and p are as defined for R¹;

R³ is hydrogen, halogen, hydroxy, C₁-6alkyl or C₁-6alkoxy;

20 R⁴ is hydrogen or C₁-6alkyl;

A is a bond or an acyclic hydrocarbon chain having 1 to 6 carbon atoms; and

n is 1 or 2.

2. A compound according to claim 1 in which R¹ is halogen, cyclohexyl, optionally substituted pyridyl or optionally substituted phenyl.

25 3. A compound according to claim 2 or 3 in which R² is hydrogen or C₁-6alkyl.

4. A compound according to any one of claims 1 to 3 in which R³ is C₁-6alkoxy.

5. A compound according to any one of claims 1 to 4 in which R⁴ is C₁-6alkyl.

30 6. A compound according to any one of claims 1 to 5 in which A is a bond.

7. A compound according to claim 1 which is:

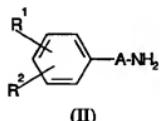
4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromo-3-methylphenyl)benzamide,

4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-bromophenyl)benzamide,

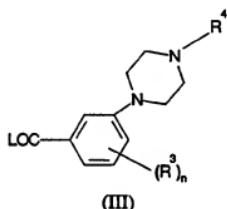
4-methoxy-3-(4-methyl-1-piperazinyl)-N-[3-methyl-4-(4-pyridyl)phenyl]benzamide,

4-methoxy-3-(4-methyl-1-piperazinyl)-N-[2'-methyl-4'-(5-methyl-1,2,4-oxadiazol-3-yl)-(1,1'-biphenyl)]benzamide, or
4-methoxy-3-(4-methyl-1-piperazinyl)-N-(4-cyclohexylphenyl)benzamide,
or pharmaceutically acceptable salts thereof.

- 5 8. A process for the preparation of a compound of formula (I) which comprises
(a) reaction of a compound of formula (II):

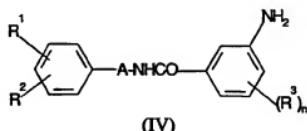


10 in which R^1 , R^2 and A are as defined in formula (I), with a compound of formula (III):



15 in which R^3 , R^4 and n are as defined in formula (I) and L is a leaving group; or

- (b) reaction of a compound of formula (IV):

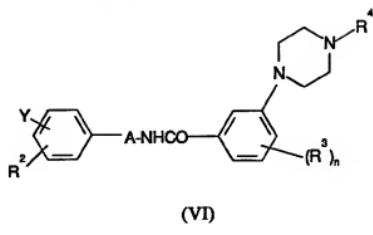


in which R¹, R², R³, A and n are as defined in formula (I) with a compound of formula (V):



in which R⁴ is as defined in formula (I) and Hal is halogen, or

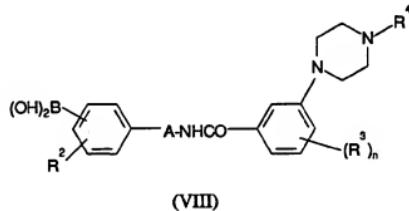
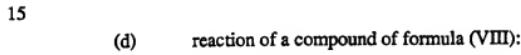
5 (c) reaction of a compound of formula (VI):



in which R², R³, R⁴, A and n are as defined in formula (I) and Y is halogen or a group -OSO₂CF₃ with a compound of formula (VII):



in which R¹ is as defined in formula (I), or

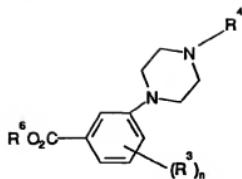


in which R², R³, R⁴, A and n are as defined in formula (I) with a compound of formula (IX):



in which R¹ is as defined in formula (I) and Y is as defined in formula (VI), or





5

in which R³, R⁴ and n are as defined in formula (I) and R⁶ is C₁₋₄alkyl in the presence of a trialkylaluminium reagent;

and optionally after any of the above processes:

- converting a compound of formula (I) into another compound of formula (I)
- 10 • forming a pharmaceutically acceptable salt.
 - 9. A compound according to any one of claims 1 to 7 for use in therapy.
 - 10. A pharmaceutical composition which comprises a compound according to any one of claims 1 to 7 in association with a pharmaceutically acceptable carrier or excipient.

15

INTERNATIONAL SEARCH REPORT

Internal Application No
PCT/EP 94/02661

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07D295/14 C07D213/40 C07D271/06 A61K31/495

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 056 144 (DR. KARL THOMAE GMBH) 21 July 1982 *Page 108-129: claims* ---	1-10
A	EP,A,0 023 569 (DR. KARL THOMAE GMBH) 11 February 1981 *Page 193-225: claims* ---	1-10
A	EP,A,0 533 268 (GLAXO GROUP LIMITED) 24 March 1993 cited in the application *Page 33-39: claims* ---	1-10
A	EP,A,0 533 267 (GLAXO GROUP LIMITED) 24 March 1993 cited in the application *Page 31-35: claims* ---	1-10
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
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- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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- "&" document member of the same patent family

Date of the actual completion of the international search

5 January 1995

Date of mailing of the international search report

13.01.95

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INTERNATIONAL SEARCH REPORT

Internal Application No
PCT/EP 94/02661

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 533 266 (GLAXO GROUP LIMITED) 24 March 1993 cited in the application *Page 24-3-93: claims* ----	1-10
P,X	JOURNAL OF MEDICINAL CHEMISTRY, vol.37, no.15, 22 July 1994 pages 2254 - 2257 JOHN W. CLITHEROW *Page 2255: table 2, compound 11; scheme 3a: step iii* *Page 2256: scheme 4a: step iii* ----	1-10

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PCT/EP 94/02661	

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